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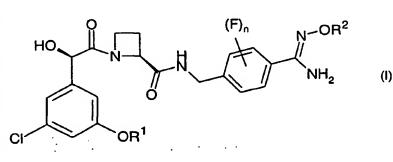
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(e.g. thrombosis) or as anticoagulants.

(57) Abstract: There is provided pharmaceutically-acceptable acid addition salts of compounds of formula (I), wherein R¹ represents C₁₋₂ alkyl substituted by one or more fluoro substituents; R² represents C₁₋₂ alkyl; and n represents 0, 1 or 2, which salts are useful as prodrugs of competitive inhibitors of trypsin-like proteases, such as thrombin, and thus, in particular, in the treatment of conditions where inhibition of thrombin is required